AMENDMENTS

What is claimed is:

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- 1. (Cancel) A method for analytical imaging of target entities, which method comprises:
 - a. obtaining a sample suspected of containing said target entities,
 - b. magnetically labeling said target entities with magnetic particles that are specific for said target entities,
 - c. magnetically manipulating said target entities towards a collection surface,
 - d. illuminating said collected target entities,
 - e. collecting sequential sub-images of said collected target entities, and
 - f. re-combining said sub-images to construct a complete image of said collected target entities.
- 2. (Cancel) The method of Claim 1, in which said target entities are cells.
- 3. (Cancel) The method of Claim 2, in which said cells are tumor cells.
- 15 4. (Cancel) The method of Claim 1, in which said magnetic labels are colloidal magnetic particles.
 - 5. (Cancel) The method of Claim 4, in which said colloidal magnetic particles are specific for the Epithelial Cell Adhesion Molecule (EpCAM).
 - (Cancel) The method of Claim 1, in which said collection surface comprises parallel Nickel lines on a glass substrate.
 - 7. (Cancel) The method of Claim 1, in which said illumination step further comprises the use of multiple wavelength light sources.
 - 8. (Cancel) An apparatus for analytical imaging of target entities, said apparatus comprising:
 - a. a sample chamber which includes a collection surface,
 - b. an arrangement of magnets capable of manipulating magnetically labeled target entities towards said collection surface.
 - c. at least one light source,
 - d. a camera capable of capturing sub-images of said collected target entities, and

- e. a computer capable of re-combining said sub-images to construct a complete image of said collected target entities.
- 9. (Cancel) The apparatus of Claim 8, in which said collection surface comprises Nickel lines on a glass substrate.
- 10. (Cancel) The apparatus of Claim 8, in which said light source is a laser.

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- 11.(Cancel) A method for automatically scanning magnetically and detectably labeled micron-sized objects located on a planar surface whereon said objects are aligned in a linear array by magnetic means, which method comprises:
 - a. loading a liquid sample containing said labeled objects into a chamber bearing a plurality of parallel magnetizable lines on said planar surface, wherein said labeled target objects have a size range of 2 to about 20 um, preferably about 5 to about 15 um;
 - b. placing said chamber on a movable magnetic x-y stage of a microscope, thereby to generate a magnetic field in proximity of said magnetizable lines, thus aligning and positionally immobilizing said objects, if present, between adjacent magnetic lines in a linear array along the x-axis;
 - c. moving said stage bearing said aligned objects along the x-axis in a digitized stepwise manner into the path of a stationary focused light beam, said light beam sequentially illuminating said aligned objects at a plurality of wavelengths each characteristic for exciting a detectable label on said target and non-target objects, thereby to generate a plurality of sequential emitted signals corresponding to segmented sub-images of said objects encoded to the specific x-y positions of the said sub-images on said stage;
 - d. acquiring and storing the sequential segmented sub-images by means of a CCD device coupled to a frame grabber at a rate commensurate with the scanning speed of the CCD device;
 - e. storing said sequential sub-images in computer memory indexed to the respective x-y-positions of said sub-images on said stage; and

- f. merging the stored sub-images of said objects to generate a reconstructed full image of each detected object, thereby to permit locating, enumerating, identifying, and classifying said objects as either target or non-target objects.
- 12. (Cancel) The method of claim 11 in which the objects are magnetically labeled by means of colloidal magnetic particles.
- 13. (Cancel) The method of claim 12 in which said colloidal magnetic particles have diameters of 50 to 300nm.
- 14. (Cancel) The method of claim 11 in which the objects are labeled with one or more detectable fluorescent substances each substantially specific for a detectable marker on said objects.
- 15. (Cancel) The method of claim 14 in which the detectable labels are selected from the groups of organic and inorganic fluorescent substances.
- 16. (Cancel) The method of claim 11 in which the objects are cells.

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- 17. (Cancel) The method of claim 11 in which said magnetic lines are about 20 to 40um wide and are separated by a distance of about 10 to 20um.
- 18. (Cancel) The method of claim 11 in which said magnetic lines are composed of a paramagnetic material.
- 19. (Cancel) The method of claim 11 in which said laser light sources have wavelengths appropriate for exciting said fluorescent substances on the labeled objects.
- 20. (Cancel) The method of claim 1 in which the CCD has a frame rate commensurate with the scan speed of the stage, thereby to maintain a resolution of at least 0.2um.

- 21. (Amended) An apparatus for automatically scanning magnetically and detectably labeled micron-sized objects on a planar surface whereon said objects are aligned in a linear array by magnetic means, comprising:
 - a. one or more laser light sources;
 - b. a polarized beam splitter with feedback detector;
 - c. a dichroic mirror assembly;

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- d. a focusing lens assembly;
- e. a sample chamber having affixed thereto at least two parallel magnetizable lines in the x-direction thereby to form a linear array, said sample chamber being inserted into a magnet system stably affixed to said x-y stage, thereby providing means for collecting, aligning and transporting said collected labeled objects into said focused light beam in a stepwise and digitized mode;
- f. means for acquiring the sequential digitized signals images emanating from said labeled objects as digitized sub-images by means of a CCD camera and one or more PMT tubes;
- g. means for storing said acquired sub-images in computer memory indexed to the corresponding z-y stage position; and
- h. means for reconstructing a full image having combined multiple digitized subimages merging said grabbed sub-images of said objects to reconstruct full images of said objects on said linear array.
- 22. (Original) The apparatus of claim 21 wherein the parallel magnetic lines on said linear array are spaced about 10 um apart.
- 23. (Original) The apparatus of claim 21 wherein the magnetic lines are composed of a paramagnetic material.
- 25 24. (Original) The apparatus of claim 21 wherein the paramagnetic material is nickel.
 - 25. (Amended) The <u>apparatus</u> method of claim 21 in which the CCD has a frame rate commensurate with the scan speed of the stage, thereby to maintain a resolution of at least 0.2um.
 - 26. (New) An apparatus for analytical imaging of target entities, said apparatus comprising:

- a. a sample chamber which includes a collection surface wherein said collection surface comprises nickel lines on a glass substrate;
- b. an arrangement of magnets capable of manipulating magnetically labeled target entities towards said collection surface;
- c. at least one light source;

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- d. a camera capable of capturing sub-images of said collected target entities; and
- e. a computer capable of re-combining said sub-images to construct a complete image of said collected target entities.
- 27. (New) The apparatus of Claim 26, in which said light source is a laser.